Asymmetric Total Synthesis of Pyranicin

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ABSTRACT

The asymmetric total synthesis of pyranicin (1) is reported. The butenolide ring was constructed via an asymmetric alkylation/ring-closing metathesis strategy. The three stereocenters in the left-hand tetrahydropyran ring were installed by sequential chiral auxiliary-mediated aldol reactions. Closure of the tetrahydropyran and fusion of the alkyl backbone were affected via a sequential ring-closing metathesis—cross-metathesis strategy.

Pyranicin (1), a novel member of the annonaceous acetogenin family of natural products, was isolated in 1997 by McLaughlin and co-workers from the stem bark of the Goniothalamus giganteus tree native to Thailand. Since 1982, over 400 molecules in the annonaceous acetogenin family have been identified, but pyranicin is one of only two known acetogenins to bear a tetrahydropyran (THP) ring. These polyether molecules in the annonaceous acetogenin family have been characterized as promising antifeedant and pesticidal treatments, as well as antimalarial, antiparasitic, and antitumor drugs, and they have recently exhibited apoptotic activity. This unique mode of biological activity has made the annonaceous acetogenins the subject of a significant amount of synthetic work. The first total synthesis of pyranicin was accomplished by Nakata and Takahashi with subsequent reports by Rein, Makabe, and Phillips.

Herein, we describe an enantioselective total synthesis of pyranicin, taking advantage of chlorotitanium enolates of N-glycoxylozazolidinones to establish the syn 1,2-oxygen relationship at C15—C16 and C19—C20. The pyranicin (PACA-2). Recent studies have further revealed in vivo cytotoxicity (ID₅₀ 9.4 µM) of pyranicin against the growth of promyelocytic leukemia cells (HL-60), alternatively attributed to its ability to inhibit DNA polymerase in the cancerous cells. The interesting structures and potent biological activity have made the annonaceous acetogenins the subject of a significant amount of synthetic work. The first total synthesis of pyranicin was accomplished by Nakata and Takahashi with subsequent reports by Rein, Makabe, and Phillips.

Herein, we describe an enantioselective total synthesis of pyranicin, taking advantage of chlorotitanium enolates of N-glycoxylozazolidinones to establish the syn 1,2-oxygen relationship at C15—C16 and C19—C20. The pyranicin

carbon backbone was envisioned to arise from a tandem ring-closing metathesis (RCM)–cross-metathesis (CM) reaction that would close the tetrahydropyran ring from triene 2 while concurrently joining the tetrahydropyran unit and butenolide fragment 3 (Figure 1). Dihydropyran precursor 2 would be accessed via an asymmetric glycolate aldol addition of glycoloyloxazolidinone 4 and aldehyde 5. The butenolide ring would be constructed via esterification of acrylic acid 7 with (S)-3-buten-2-ol (6), followed by RCM.

Aldehyde 5 was prepared from (S)-benzylglycidyl ether as illustrated in Scheme 1. Lewis acid promoted addition of lithiated homopropargyl alcohol 8 to (S)-benzyl glycidyl ether provided alkyne 9. The alkyne was reduced, and removal of the benzyl group was accomplished employing Raney nickel to deliver diol 10. Selective sulfonylation of the primary alcohol was best affected employing 2,4,6-trisopropylsulfonyl chloride (TrisCl) under standard conditions whereupon treatment with base afforded epoxide 11. Subsequently, the (S)-epoxide underwent copper(I)-promoted reaction with butenylmagnesium bromide to provide alcohol 12. Ensuing alcohol protection, selective removal of the PMB ether, and Swern oxidation of the primary alcohol provided the target aldehyde 5 in good yield over three steps.

Preparation of triene 2 began with a glycolate aldol reaction between benzylglycoloyloxazolidinone 13 and tridecanal, providing aldol adduct 14 in good yield and excellent diastereoselectivity (Scheme 1). This reaction established the stereocenters at C19 and C20 at an early stage. The secondary alcohol was then protected as its triethylsilyl (TES)
ether, and the chiral auxiliary was reductively removed with lithium borohydride.

Oxidation\textsuperscript{13} of primary alcohol \textsuperscript{15} followed by Wittig methylenation provided the protected diol, which was selectively deprotected under fluoride conditions to give secondary alcohol \textsuperscript{16}. Subsequent alkylation of the free alcohol with bromoacetic acid gave the glycolic acid, and further transformation into glycolylimide \textsuperscript{4} was accomplished via nucleophilic addition of lithiated oxazolidinone \textsuperscript{17} to the intermediate mixed pivaloyl anhydride.\textsuperscript{14}

A second titanium-mediated glycolate aldol reaction\textsuperscript{10} with aldehyde \textsuperscript{5} established the stereocenters at C15 and C16 providing the aldol adduct \textsuperscript{18} in 74% yield (\textsuperscript{95}:5 diastereoselectivity). The RCM precursor \textsuperscript{2} was prepared from aldol adduct \textsuperscript{18} by a four-step sequence. Protection of the C15 hydroxyl as its TES ether followed by reductive removal of the auxiliary gave the primary alcohol \textsuperscript{19}. Dess–Martin oxidation\textsuperscript{15} of the alcohol to the aldehyde and final methylenation completed the synthesis of triene \textsuperscript{2}.

Our efforts were directed next toward the preparation of \(\gamma\)-methylbutenolide \textsuperscript{3}. The C34 stereocenter was to be installed via esterification using (S)-3-buten-2-ol (6). Although the enantiomer of the alcohol had previously been prepared in our total synthesis of giganticin,\textsuperscript{16} the volatility of the alcohol (bp 92°C) created difficulty with its successful isolation. In an attempt to ease the problems with isolation of alcohol 6, while still maintaining the necessary terminal olefin functionality for the ensuing RCM reaction, we investigated the use of alcohol \textsuperscript{22} in an alternative relay ring-closing metathesis (RCRM) strategy, recently demonstrated by Hoye\textsuperscript{17} (Scheme 2). We anticipated that while the increased molecular weight of the ether fragment of alcohol \textsuperscript{22} would effectively improve the isolation of the chiral alcohol, the additional atoms would be removed as dihydrous butenolide \textsuperscript{3}. The synthesis of bis-allylic ether \textsuperscript{22} was thus pursued. L-Ethyl lactate was protected as its tert-butyldiphenyldisilyl (TBDPS) ether \textsuperscript{20}, which was subsequently reduced to the corresponding aldehyde (Scheme 2). Olefination of the intermediate aldehyde employing carboethoxymethylene triphenylphosphorane yielded \(R,\beta\)-unsaturated ester \textsuperscript{21}, which was then reduced to the allylic alcohol. Alkylation of the alcohol with allyl bromide gave the allyl ether, which was exposed to \(n\)-Bu\textsubscript{4}NF to provide the desired alcohol \textsuperscript{22} in 80% over two steps.

The synthesis of acrylate \textsuperscript{7} began with installation of the C4 stereocenter via asymmetric alkylation\textsuperscript{18} of \(p\)-methoxybenzylglycolyloxazolidinone \textsuperscript{23} with allylic iodide \textsuperscript{24},\textsuperscript{19} giving protected allyl alcohol \textsuperscript{25} with high (>95:5) diastereoselectivity (Scheme 2). Reductive removal of the chiral auxiliary unmasked a primary alcohol \textsuperscript{26}, which allowed for

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conversion to the terminal alkene via Swern oxidation and subsequent olefination with methyleneetriphenylphsophorane. Removal of the primary TBS ether followed by a two-stage oxidation yielded the corresponding acrylic acid. Esterification of acid with alcohol was best affected via the intermediate mixed pivaloyl anhydride. Initial attempts at closure of the butenolide under standard RRCM conditions only gave the corresponding ring-opened diene, presumably formed after the initial expulsion of dihydrofuran, followed by intermolecular carbene transfer. However, by sparging the reaction with argon, we were able to increase the yield of butenolide from 23% to 70%, without any trace of straight-chain byproduct.

The completion of the synthesis required the closure of the tetrahydropyran ring via a ring-closing metathesis and assembly of the THP and butenolide fragments through a cross-metathesis reaction (Scheme 3). Given the expected rapid closure of the tetrahydropyran and the difference in reactivity between the C5 alkene of the butenolide and the C6 alkene of the tetrahydropyran unit, it was hoped that the two processes could be accomplished concurrently in a single exposure to a ruthenium carbene.

Despite the differences in the reactivities of the two coupling fragments, however, the ensuing tandem RCM/CM reaction proved to be quite difficult to control, and suitable conditions for the tandem sequence could not be identified. The best results were found when the cross-metathesis was performed between the ring-closed monomer and butenolide. Therefore, it was necessary to isolate the ring-closed monomer before subjecting it to cross-metathesis with butenolide. In the event, triene was exposed to the Grubbs second-generation catalyst to close the tetrahydropyran ring. After the ring-closing metathesis product was isolated in quantitative yield, the cross-metathesis between alkene and butenolide was achieved with the Hoveyda–Grubbs catalyst in 58% yield based on recovered starting material.

Finally, exposure of triene to excess tosylhydrazine and sodium acetate in aqueous dimethoxyethane at reflux resulted in selective hydrogenation of the C17–C18 and C5–C6 alkenes and removal of the triethylsilyl protecting group, while leaving the butenolide olefin in tact. The synthetic sample was identical in all aspects (1H, 13C, [α]D) to the natural product.

In summary, a highly convergent total synthesis of pyranicin has been completed. The key fragments were constructed employing asymmetric glycolate aldol and alkylation reactions followed by ring-closing metatheses; a cross-metathesis was utilized to convergently construct the backbone of the natural product.

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Supporting Information Available: Experimental procedures and copies of 1H and 13C NMR spectra. This material is available free of charge via the Internet at http://pubs.acs.org.

Scheme 3. Completion of the Synthesis


